

STIC-ILL

vol w 437391

**From:** STIC-Biotech/ChemLib  
**Sent:** Thursday, March 20, 2003 7:20 AM  
**To:** STIC-ILL  
**Subject:** FW: 09850697

-----Original Message-----

**From:** Yaen, Christopher  
**Sent:** Wednesday, March 19, 2003 5:00 PM  
**T :** STIC-Biotech/ChemLib  
**Subject:** 09850697

10027135

could you please get the following ref(s):

Thromb Haemost 1989 Nov 24;62(3):846-9

2595658

Appl Environ Microbiol 1994 Aug;60(8):2793-801

Antonie Van Leeuwenhoek 1996 Feb;69(2):151-9

Oncogene 2000 Mar 16;19(12):1579-88

Christopher Yaen  
Patent Examiner  
US PTO  
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CM1-Rm 8E18  
Mail Box 8E12  
703-305-3586

STIC-ILL

NO

437370

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**From:** Yaen, Christopher  
**Sent:** Wednesday, March 19, 2003 5:00 PM  
**To:** STIC-Biotech/ChemLib  
**Subject:** 09850697

10026554

could you please get the following ref(s):

Thromb Haemost 1989 Nov 24;62(3):846-9

Appl Environ Microbiol 1994 Aug;60(8):2793-801

Antonie Van Leeuwenhoek 1996 Feb;69(2):151-9

8775975

Oncogene 2000 Mar 16;19(12):1579-88

Christopher Yaen  
Patent Examiner  
US PTO  
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NEWS 26 OCT 01 EVENTLINE has been reloaded  
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NEWS 39 JAN 13 NUTRACEUT offering one free connect hour in February 2003  
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NEWS 41 JAN 21 Simultaneous left and right truncation added to COMPENDEX,  
ENERGY, INSPEC  
NEWS 42 JAN 29 CANCERLIT is no longer being updated  
NEWS 43 FEB 13 METADEX enhancements  
NEWS 44 FEB 24 PCTGEN now available on STN  
NEWS 45 FEB 24 TEMA now available on STN  
NEWS 46

NEWS 47 Feb 26 NTIS now allows simultaneous left and right truncation  
NEWS 48 Feb 26 PCTFULL now contains images  
NEWS 49 Mar 04 SDI PACKAGE for monthly delivery of multifile SDI results  
NEWS 50 Mar 19 APOLLIT offering free connect time in April 2003

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=> S SPAF 202 SPAF

L1

==> s l1 and vector  
 L2 13 L1 AND VECTOR  
  
 ==> dup rem l2  
 PROCESSING COMPLETED FOR L2  
 L3 12 DUP REM L2 (1 DUPLICATE REMOVED)  
  
 ==> d 1-12  
  
 L3 ANSWER 1 OF 12 PCTFULL COPYRIGHT 2003 Univentio  
 AN 2003014322 PCTFULL ED 20030303 EW 200308  
 TITN PROTEINS ASSOCIATED WITH CELL GROWTH, DIFFERENTIATION, AND DEATH  
 TIFR PROTEINES ASSOCIEES A LA CROISSANCE, LA DIFFERENTIATION ET LA MORT  
 CELLULAIRES  
  
 IN AZIMZAI, Yalda, 5518 Boulder Canyon Drive, Castro Valley, CA 94552, US  
 (US, US);  
 BARROSO, Ines, 38 Eden Street, Cambridge, Kent CB1 1EL, GB [PT, GB];  
 BAUGHN, Mariah, R., 14244 Santiago Road, San Leandro, CA 94577, US [US,  
 US];  
 BECHA, Shanya, D., 21062 Gary Drive # 117, Castro Valley, CA 94546, US  
 (US, US);  
 BOROWSKI, Mark, L., 122 Orchard Avenue, Redwood City, CA 94061, US [US,  
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 DUGGAN, Brendan, M., 243 Buena Vista Avenue # 306, Sunnyvale, CA 94086,  
 US [AU, US];  
 ELLIOTT, Vicki, S., 3770 Polton Place Way, San Jose, CA 95121, US [US,  
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 US], for US only;  
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 for US only;  
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 YUE, Henry, 826 Lois Avenue, Sunnyvale, CA 94087, US [US, US];  
 YUE, Huibin, 1170 South Stelling Road, Cupertino, CA 95014, US [US, US]

AG

CA 94304, US  
English  
LA  
English  
DT  
Patent  
PI  
WO 2003014322 A2 20030220  
W: AE AG AL AM AT AU AZ BA BB BG BR BY BZ CA CH CN CO CR CU CZ  
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KE KG KP KR KZ LC LK LR LS LT LU LV MA MD MG MK MN MW MX NZ  
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RW (ARIPO): GH GM KE LS MW NZ SD SL SZ TZ UG ZM ZW  
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RW (OAPI): BF BJ CF CG CI CM CA GN GQ GW ML MR NE SN TD TG  
AI WO 2002-US25465 A 20020808  
PRAI US 2001-60/311.017 20010808  
US 2001-60/313.070 20010817  
US 2001-60/313.071 20010817  
US 2001-60/314.678 20010824  
US 2001-60/316.692 20010831  
US 2001-60/317.913 20010907  
US 2001-60/322.182 20010914  
US 2001-60/340.747 20011207  
US 2001-60/342.761 20011220  
US 2002-60/369.129 20020329  
L3 ANSWER 2 OF 12 CAPLUS COPYRIGHT 2003 ACS DUPLICATE 1  
AN 2002:755245 CAPLUS  
DN 137:274175  
TI DNA, CDNA and protein sequences of spermatogenesis assocd. factors a  
method for diagnosis of cancer  
IN Kulcsz-Martin, Molly F.; Liu, Yungang  
PA USA  
U.S. Pat. Appl. Publ.. 42 pp., Cont.-in-part of U.S. Ser. No. 777,753.  
CODEN: USXXCO  
DT Patent  
LA English  
FAN CNT 1  
PATENT NO. KIND DATE APPLICATION NO. DATE  
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PI US 2002143169 A1 20021003 US 2001-850697 20010508  
PRAI US 1997-938708 B2 19970927  
US 2001-777753 A2 20010206  
L3 ANSWER 3 OF 12 USPTATFULL  
AN 2002:301167 USPTATFULL  
TI Nucleic acids, proteins, and antibodies  
IN Rosen, Craig A., Laytonville, MD, UNITED STATES  
Ruben, Steven N., Olney, MD, UNITED STATES  
Barash, Steven C., Rockville, MD, UNITED STATES  
PI US 2002168711 A1 20021114  
AI US 2001-764868 A1 20010117 (9)  
PRAI US 2000-179065P 20000131 (60)  
US 2000-180628P 20000204 (60)  
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US 2000-237040P 20001002 (60)  
US 2000-240960P 20001020 (60)  
US 2000-239935P 20001013 (60)  
DT Utility  
FS APPLICATION  
LN CNT 31967  
INCL INCLM: 435/069.100  
INCL: 435/325.000; 435/320.100; 435/183.000; 530/350.000; 536/023.100  
NCL INCLM: 435/069.100  
NCL: 435/325.000; 435/320.100; 435/183.000; 530/350.000; 536/023.100  
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ICM: C12P021-02  
ICS: C12N005-06; C07H021-04; C12N009-00; C07K014-435  
CAS INDEXING IS AVAILABLE FOR THIS PATENT.  
L3 ANSWER 4 OF 12 USPTATFULL  
AN 2002:112541 USPTATFULL  
TI Proteins related to schizophrenia and uses thereof  
IN St. George-Hyslop, Peter H., Toronto, CANADA  
Fraser, Paul E., Toronto, CANADA  
PA The Governing Council of the University of Toronto (non-U.S. corporation)  
PI US 2002058276 A1 20020516  
AI US 2001-945258 A1 20010831 (9)  
PRAI US 2000-229889P 20000901 (60)  
DT Utility  
FS APPLICATION  
LN CNT 2909  
INCL INCLM: 435/006.000  
INCL: 424/009.200; 800/003.000  
NCL INCLM: 435/006.000

IC NCLS: 424/009.200; 800/003.000

AN ICM: C12Q001-68

ICS: A61K049-00; A01K067-00

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L3 ANSWER 5 OF 12 PCTFULL COPYRIGHT 2003 Univentio  
AN 2002099062 PCTFULL ED 20021218 EW 200250  
TIEN NOVEL ANTIBODIES THAT BIND TO ANTIGENIC POLYPEPTIDES, NUCLEIC ACIDS  
ENCODING THE ANTIGENS, AND METHODS OF USE  
TIFR NOUVEAUX ANTICORPS SE FIXANT A DES POLYPEPTIDES ANTIGENIQUES, ACIDES  
NUCLEIQUES CODANT LES ANTIGENES ET MODES D'UTILISATION  
IN ANDERSON, David, W., 85 Montoya Drive, Branford, CT 06405, US [US];  
ZERHUSEN, Bryan, D., 337 Monticello Drive, Branford, CT 06405, US [US];

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06511, US [US, US], for all designates States except US;  
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English  
LA English  
DT Patent  
PI WO 2002099062  
DS W: A2 20021212

AE AG AL AM AT AU AZ BA BB BG BR BY BZ CA CH CN CO CR CU CZ  
DE DK DM DZ EE ES ET GB GD GE GH GM GR GU HR HU ID IL IN IS JP  
KE KG KP KR KZ LC LK LR LS LT LU LV MA MD MG MK MN MW MX MZ  
NO NZ OM PH PL PT RO RU SD SE SG SI SK SL TJ TM TN TR TT T2

UA UG US UZ VN YU ZA ZM ZW  
RW (ARIPO): GH GM KE LS MW MZ SD SL SZ TZ UG ZM ZW  
RW (EAP): AM AZ BY KG KZ MD RU TJ TM

RW (EPO): AT BE CH CY DE DK ES FI FR GB GR IE IT LU MC NL PT SE TR

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           GRIFFIN, Jennifer, A., 33691 Mello Way, Fremont, CA 94555, US [US, US];            YUE, Henry, 826 Lois Avenue, Sunnyvale, CA 94087, US [US, US];            LEE, Ernestine, A., 624 Kains Street, Albany, CA 94706, US [US, US];            BAUGHN, Mariah, R., 14244 Santiago Road, San Leandro, CA 94577, US [US, US];            DUGGAN, Brendan, M., 243 Buena Vista Avenue #306, Sunnyvale, CA 94086, US [AU, US];            WALIA, Narinder, K., 890 Davis Street, #205, San Leandro, CA 94577, US [US, US];            LEE, Sally, 825 East Evelyn, #425, Sunnyvale, CA 94086, US [US, US];            RAMKUMAR, Jayalaxmi, 34359 Maybird Circle, Fremont, CA 94555, US [IN, US];            WARREN, Bridget, A., 10130 Parkwood Drive #2, Cupertino, CA 95014, US [US, US];            GANDHI, Ameena, R., 705 5th Avenue, San Francisco, CA 94118, US [US, US];            LU, Dyung, Aina, M., 233 Coy Drive, San Jose, CA 95123, US [US, US];            LU, Yan, 3885 Corrina Way, Palo Alto, CA 94303, US [CN, US];            YAO, Monique, G., 1189 Woodgate Drive, Carmel, IN 46033, US [US, US];            DING, Li, 3353 Alma Street #146, Palo Alto, CA 94306, US [CN, US];            TRIBOULEY, Catherine, M., 1121 Tennessee Street, #5, San Francisco, CA 94107, US [FR, US];            SANJANWALA, Madhu, M., 210 Sylvia Court, Los Altos, CA 94024, US [US, US];            ARVIZU, Chandra, 490 Sherwood Way #1, Menlo Park, CA 94025, US [US, US];            HILLMAN, Jennifer, L., 230 Monroe Drive, #17, Mountain View, CA 94040, US [US, US];            HAMLET-COX, Diana, Incyte Genomics, Inc., 3160 Porter Drive, Palo Alto, CA 94304, US [English            Patent            WO 2002046385            W: AE AG AL AM AT AU AZ BA BB BG BR BY BZ CA CH CN CO CR CU CZ DE DK DM DZ EC EE ES FI GB GD GE GH GM HR HU ID IL IN IS JP KE KG KP KR KZ LC LK LR LS LT LU LV MA MD MG MN MW MX MZ NO NZ PL PT RO RU SD SE SG SI SK SL TJ TM TR TT TZ UA UG US UZ VN YU ZA ZW            RW (ARIPO): GH GM KE LS MW MZ SD SL SZ TZ UG ZM ZW            RW (EAP): AM AZ BY KG KZ MD RU TJ TM            RW (EPO): AT BE CH CY DE DK ES FI FR GB GR IE IT LU MC NL PT SE TR            RW (OAPI): BF BJ CF CG CI CM GA GN GQ GW ML MR NE SN TD TG            WO 2001-US47432 A 20011204            US 2000-60/251,824 20001207            US 2000-60/254,312 20001208            US 2000-60/255,773 20001214            US 2000-60/256,188 20001215            US 2000-60/255,940 20001215            US 2000-60/257,488 20001221            US 2001-60/262,839 20010119            US 2001-60/264,402 20010126            CL2N009-00         </p>	<p>           AG            LAF            LA            English            Patent            WO 2002046385            W: AE AG AL AM AT AU AZ BA BB BG BR BY BZ CA CH CN CO CR CU CZ DE DK DM DZ EC EE ES FI GB GD GE GH GM HR HU ID IL IN IS JP KE KG KP KR KZ LC LK LR LS LT LU LV MA MD MG MN MW MX MZ NO NZ PL PT RO RU SD SE SG SI SK SL TJ TM TR TT TZ UA UG US UZ VN YU ZA ZW            RW (ARIPO): GH GM KE LS MW MZ SD SL SZ TZ UG ZM ZW            RW (EAP): AM AZ BY KG KZ MD RU TJ TM            RW (EPO): AT BE CH CY DE DK ES FI FR GB GR IE IT LU MC NL PT SE TR            RW (OAPI): BF BJ CF CG CI CM GA GN GQ GW ML MR NE SN TD TG            WO 2001-US47432 A 20011204            US 2000-60/251,824 20001207            US 2000-60/254,312 20001208            US 2000-60/255,773 20001214            US 2000-60/256,188 20001215            US 2000-60/255,940 20001215            US 2000-60/257,488 20001221            US 2001-60/262,839 20010119            US 2001-60/264,402 20010126            CL2N009-00         </p>	<p>           AI            PRAI         </p>	<p>           ICM         </p>
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US 2000-60/249, 265 20001117  
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US 2000-60/250, 160 20001201  
US 2000-60/256, 719 20001205  
US 2000-60/251, 030 20001205  
US 2000-60/251, 988 20001205  
US 2000-60/251, 479 20001206  
US 2000-60/251, 869 20001208  
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US 2000-60/251, 868 20001208  
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ANSWER 11 OF 12 PCTFULL COPYRIGHT 2003 Univentio  
2001045007 PCTFULL ED 20020827  
A METHOD AND SYSTEM FOR DISCOVERY OF TRADES BETWEEN PARTIES  
PROCEDE ET SYSTEME PERMETTANT DE DECOUVRIR DES MARCHES ENTRE DES PARTIES  
MACREADY, William, G.;  
EL-BELTAGY, Mohammed;  
ROY, Barbeau;  
ANDERSON, Mark  
BIOS GROUP INC.  
Patent

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AE AG AL AM AT AU AZ BA BB BG BR BY BZ CA CH CN CR CU CZ DE  
DK DM DZ EE ES FI GB GD GE GH GM GR HU ID IL IN IS JP KE KG  
KR KZ LC LR LS LT LU LV MA MD MG MK MN MX MY NZ NO NZ  
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RU TJ TM AT BE CH CY DE DK ES FI FR GB GR IE IT LU MC NL PT  
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ANSWER 12 OF 12 PCTFULL COPYRIGHT 2003 Univentio  
2000060069 PCTFULL ED 20020515  
A PRESENILIN ASSOCIATED MEMBRANE PROTEIN AND USES THEREOF  
PROTEINE MEMBRANAIRE ASSOCIEE A LA PRESENILINE ET SES UTILISATIONS  
ST. GEORGE-HYSLOP, Peter, H.;  
FRASER, Paul, E.  
THE GOVERNING COUNCIL OF THE UNIVERSITY OF TORONTO  
English  
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AE AG AL AM AT AU AZ BA BB BG BR BY CA CH CN CR CU CZ DE DK  
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GH GM KE LS MW SD SL SZ TJ ZY ZY ZY ZY ZY ZY ZY ZY ZY ZY  
AT BE CH CY DE DK ES FI FR GB GR IE IT LU MC NL PT SE BF BJ  
CF CG CI CM CA CN CH ML MR NE SN TD TG

WO 2000-CA354  
US 1999-60/127, 452 19990401  
US 1999-60/173, 826 19991230  
C12N015-12  
C07K014-705; A01K067-027; C12N005-10; C12Q001-68; G01N033-50

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L3 ANSWER 12 OF 12 PCTFULL COPYRIGHT 2003 Univentio

DETD Furthermore, mutant PAMP nucleic acids, proteins, or peptides, cells transfected with vectors comprising mutant PAMP nucleic acids, transgenic animals expressing mutant PAMP or peptides thereof, and their use in studying Alzheimer's Disease and other.

The invention also provides vectors, and particularly expression vectors (e.g., cos-Tet vector), which include any of the above-described SUBSTITUTE SHEET (RULE 26) nucleic acids. It is a further object of the invention to provide vectors in which normal or mutant PAMP nucleic acid sequences are operably joined to exogenous regulatory regions to produce altered patterns of expression.

phenotype on the cell in which it is expressed. The term expression system means a host cell transformed by a compatible expression vector and cultured under suitable conditions e.g. for the expression of a protein coded for by foreign DNA carried by the vector and introduced to the host cell.

The terms vector, cloning vector and expression vector mean the vehicle by which a DNA or RNA sequence (e.g., a foreign gene) can be introduced into a host cell, so.

Vectors include plasmids, phages, viruses, etc. A cassette refers to a DNA coding sequence or segment of DNA that codes for an expression product that can be inserted into a vector at defined restriction sites. The cassette restriction sites are designed to ensure insertion of the cassette in the proper reading frame. Generally, foreign DNA is inserted at one or more sites of the vector DNA, and then is carried by the vector into a host cell along with the transmissible vector DNA. A segment or sequence of DNA having inserted or added DNA, such as an expression vector, can also be called a DNA construct. Recombinant cloning vectors will often include one or more SUBSTITUTE SHEET (RULE 26) replication systems for cloning or expression, one or more markers for selection in.

In the context of the present invention, an gene is heterologous to the recombinant vector DNA in which it is inserted for cloning or expression, and it is heterologous to a host cell containing such a vector, in which it is expressed, e.g., a CHO cell.

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pumps inserted into the cerebral ventricles); via the transplantation of genetically-modified cells expressing recombinant genes; or via the use of biological vectors (e.g., retrovirus, adenovirus, adeno-associated virus, lentivirus, or herpes simplex virus-based vectors) which allow expression of appropriately modified gene products in selected cell types. It should be noted that the recombinant proteins described. . . or part of PAMP or PAMP mutants, e.g., as mini-gene cDNA transgene constructs under the regulation of suitable promoter elements carried in vectors such as cos-Tet for transgenic mice and pcDNA (Invitrogen, California) in transfected cell lines.

to identify upstream and downstream modifiers of a PAMP phenotype. Transgenic animals can also be prepared by introducing the transgene on a vector: such animals, which are not modified in the germ line and are only transiently, naturally cannot pass along the genetic information. . . may be truncated), 1.5 can be introduced in vivo, ex vivo, or in vitro using a viral or a non-viral vector, e.g., as discussed above. Expression in targeted tissues can be effected by targeting the transgenic vector to specific cells, such as with a viral vector or a receptor ligand, or by using a tissue-specific promoter, or both. Targeted gene delivery is described in International Patent Publication WO. . . Preferably, for in vivo administration, an appropriate immunosuppressive treatment is employed in conjunction with the viral vector.

e.g., adenovirus vector, to avoid immuno-deactivation of the viral vector and transduced cells. For example, immunosuppressive cytokines, such as interleukin 12 (IL-12), interferon- $\gamma$  (IFN $\gamma$ ), or anti-CD4 antibody, can be administered to block humoral or cellular immune responses to the viral vectors (see, e.g., Wilson, Nature Medicine, 1995). In that regard, it is advantageous to employ a viral vector that is engineered to express a minimal number of antigens.

Herpes virus vectors. Because herpes virus is tropic for cells of the nervous system (neural cells), it is an attractive vector for delivery of function PAMP genes. Various defective (non-replicating, and thus non-infectious) herpes virus vectors have been described, such as a defective herpes virus 1 (HSV1) vector (Kaplit et al., Molec. Cell. Neurosci. 2:320-330, 1991, International Patent Publication No. WO 94/21807, published September 29, 1994; International. . .

Adenovirus vectors. Adenoviruses are eukaryotic DNA viruses that can be modified to efficiently deliver a nucleic acid of the invention to a . . . adenovirus, more preferably a CAV2 adenovirus (e.g., Mahattan or A26/61 strain (ATCC VR-800), for example). Various replication defective adenovirus and minimum adenovirus vectors have been described for gene therapy (WO94126914, WO95/026971 WO94/28938, SUBSTITUTE SHEET (RULE 26) which is infected with a human helper virus (for. . .

Retrovirus vectors. In another embodiment the gene can be introduced in a retroviral vector, e.g., as described in Anderson et al., U.S.

infect dividing cells. The retrovirus genome includes two LTRs, an encapsidation sequence and three coding regions (gag, pol and env). In recombinant retroviral vectors, the gag, pol and env genes are generally deleted, in whole or in part, and replaced with a heterologous nucleic acid. . . These vectors can be constructed from different types of retrovirus, such as MoMuLV (murine Moloney leukemia virus), MEV (murine Moloney sarcoma virus), HaSV (Harvey. . . line PA317 (US 4,861,719); the PsiCRIP cell line (WO 90/02806) and the GP-envAm-12 cell line (WO 89/07150). In addition, the recombinant retroviral vectors can contain modifications within the LTRs for suppressing transcriptional activity as well as extensive encapsidation sequences which may include a part of the gag gene (Bender et al., J. Virol. 61:1639, 1987). Recombinant retroviral vectors are purified by standard techniques known to those having ordinary skill in the art.

Retrovirus vectors can also be introduced by recombinant DNA viruses, which permits one cycle of retroviral replication and amplifies transfection efficiency (see WO 95/22617. . .

Lentivirus vectors. In another embodiment, lentiviral vectors are can be used as agents for the direct delivery and sustained expression of a transgene in several tissue types, including brain, retina, muscle, liver and blood. The vectors can efficiently transduce dividing and non-dividing cells in . . . SUBSTITUTE SHEET (RULE 26) these tissues, and maintain long-term expression of the gene of. . . 72,9873-801,1998). Lentiviral packaging cell lines are available and known generally in the art. They facilitate the production of high-titer lentivirus vectors for gene therapy. An example is a tetracycline-inducible VSV-G pseudotyped lentivirus packaging cell line which can generate virus particles at titers greater than 10<sup>6</sup> IU/ml for at least 3 to 4 days (Kafri, et al., J. Virol., 73: 576-584, 1999). The vector

produced by the inducible cell line can be concentrated as needed for efficiently transducing nondividing cells in vitro and in vivo.

Non-viral vectors. A vector can be introduced in vivo in a non-viral vector, e.g., by lipofection, with other transfection facilitating agents (peptides, polymers, etc.), or as naked DNA. Synthetic cationic lipids can be used to . . .

DNA vectors for gene therapy can be introduced into the desired host cells by methods known in the art, e.g., electroporation, microinjection, cell fusion, DEAE dextran, calcium phosphate precipitation, use of a gene gun (ballistic transfection), or use of a DNA vector transporter (see, e.g., Wu et al., J. Biol. . .

either V5-tagged PAMP or empty plasmid (mock transfection control). Duplicate experiments were performed by: (1) transient transfection of V5-PAMP and PAPP695 (or empty vector plus PAPP695 as a mock transfection control) into murine embryonic fibroblasts stably infected with human PSI expressed from a retroviral vector construct (Clontech, CA) - or (2) transient transfection of V5-PAMP (or empty plasmid) into HEK293 cell lines with a stable expression of . . .

Cells were transiently transfected with PAMP cDNA (SEQ ID NO: 13) tagged at the 3'-end with a V5-epitope encoded from the pCDNA6 vector. The conditioned media were collected 20 hr after transient transfection with PAMP (or with empty vector), and the Ap40 and Ap42 levels were measured by ELISA (Zhang L, et al., J Biol Chem 1999; 274: 8966 ). In . . .

region D369L PAMR0312-369 in the central conserved region D340X: PAMPA312-340 in the central conserved region YDT-- PAMPD458A in the putative 'aspartyl protease' DTA site SPAP- PAMPP633A/F635A in the SPAP motif TM: PAMPS683A in the TM domain C31D: PAMPA630-668 in the conserved region adjacent to the TM domain To further examine the role of . . . above mutations, as well as normal/wild type PAMP (PAMPwt) ONA and the ONA for an unrelated protein (LacZ), in frame into pCDNA6 vectors. A series of HEK293 cell lines stably expressing endogenous PSI 7 PAPPswedish and either wild type PAMP or PAMP constructs in which. . .

or in the AP42/AK ratio, when the PAMPwt, PAMPD458A, PAMPA630-668, PAMPP633A/F635A, and PAMPS683A cells were compared to control lines (expressing LacZ, or . . .

empty vector).

Mock (LacZ/empty 1.0 1.0 1.0 vector)

Wild type nicastrin 1.03  $\pm$ plusmn; 0.09 1.05  $\pm$ plusmn; 0.07 0.99  $\pm$ plusmn; 0.07 0.07 0.07 D36A/V337A 3.09  $\pm$ plusmn; 0.50 1.61  $\pm$ plusmn; 0.19 1.81  $\pm$ plusmn; 0.15 0.15 0.15 (p . . .

CLMEN 10 A vector comprising the nucleic acid of claim 7 operatively associated with an expression control sequence.

11 A cell transfected with the vector of claim 1 0.

15 A vector comprising the nucleic acid of claim 13 operatively associated with an expression control sequence.

0 16. A cell transfected with the vector of claim 15.

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L3 ANSWER 1 OF 12 FCTFULL COPYRIGHT 2003 Univentio ABEN . . . cell growth, differentiation, and death (CGDD) and polynucleotides which identify and encode CGDD. Embodiments of the invention also provide expression vectors, host cells, antibodies, agonists, and antagonists. Other embodiments provide methods for diagnosing, treating, or preventing disorders associated with aberrant expression. . .

DETD . . . Hamada, K. et al. (1996; Cancer Res. 56:3047-3054) are investigating the introduction of p53 into cervical cancer cells via an adenoviral vector as an experimental therapy for cervical cancer.

Spermatogenesis associated factor (SPAF) is an AAA-protein (Afpase associated with diverse activities) specific to early spermatogenesis and malignant conversion. SPAF is expressed in spermatogonia and early spermatocytes in the basal compartment of the seminiferous tubules (Liu, Y.

Table 6 provides an appendix which describes the tissues and vectors used for construction of the cDNA libraries shown in Table 5.

are now described. All publications mentioned herein are cited for the purpose of describing and disclosing the cell lines, protocols, reagents and vectors which are reported in the publications and which might be used in connection with various embodiments of the invention. Nothing herein. . .

include a nucleic acid sequence operably linked to a promoter sequence. Such a recombinant nucleic acid may be part of a vector that is used, for example, to transform a cell.

Alternatively, such recombinant nucleic acids may be part of a viral vector, e.g., based on a vaccinia virus, that could be used to vaccinate a mammal wherein the recombinant nucleic acid is expressed, inducing. . .

or by infection with a recombinant virus. In another embodiment, the nucleic acid can be introduced by infection with a recombinant viral vector, such as a lentiviral vector (Lois, C. et al. (2002) Science 295:868-872). The term genetic manipulation does not include classical cross-breeding, or in vitro fertilization, but rather.

is 53% identical from residue M1 to residue K56, and 84% identical from residue G55 to residue G535, to *Mus musculus* SPAP (spermatogenesis associated factor, AAA family) (GenBank ID g4105619) as determined by the Basic Local Alignment Search Tool (BLAST). (See Table 2.) The.

most frequently represented by the Incyte cDNA sequences which were used to assemble and confirm the above polynucleotides. The tissues and vectors which were used to construct the cDNA libraries shown in Table 5 are described in Table 6. fragments thereof, entirely by synthetic chemistry. After production, the synthetic polynucleotide may be inserted into any of the many available expression vectors and cell systems using reagents well known in the art. Moreover, synthetic chemistry may be used to introduce mutations into a polynucleotide.

method which may be employed, restriction-site PCR, uses universal and nested primers to amplify unknown sequence from genomic DNA within a cloning vector (Sarkar, G. (1993) PCR Methods Appl. 2:318-322). Another method, inverse PCR, uses primers that extend in divergent directions to amplify unknown.

order to express a biologically active CGDD, the polynucleotides encoding CGDD or derivatives thereof may be inserted into an appropriate expression vector, i.e., a vector which contains the necessary elements for transcriptional and translational control of the inserted coding sequence in a suitable host. These elements include regulatory sequences, such as enhancers, constitutive and inducible promoters, and 5' and 3' untranslated regions in the vector and in polynucleotides encoding CGDD. Such elements may vary in their strength and specificity. Specific initiation signals may also be used to. . . in cases where a polynucleotide sequence encoding CGDD and its initiation codon and upstream regulatory sequences are inserted into the appropriate expression vector, no additional transcriptional or translational control signals may be needed. However, in cases where only coding sequence, or a fragment thereof, is inserted, exogenous translational control signals including an in-frame ATG initiation codon should be provided by the vector. Exogenous translational elements and initiation codons may be of various origins, both natural and synthetic. The efficiency of expression may be.

Methods which are well known to those skilled in the art may be used to

construct expression  
72

vectors containing polynucleotides encoding CGDD and appropriate transcriptional and translational control elements. These methods include in vitro recombinant DNA techniques, synthetic techniques, and.

A variety of expression vector/host systems may be utilized to contain and express polynucleotides encoding CGDD. These include, but are not limited to, microorganisms such as bacteria transformed with recombinant bacteriophage, plasmid, or cosmid DNA expression vectors; yeast transformed with yeast expression vectors; insect cell systems infected with viral expression vectors (e.g., baculovirus); plant cell systems transformed with viral expression vectors (e.g., cauliflower mosaic virus, CaMV, or tobacco mosaic virus, TMV) or with bacterial expression vectors (e.g., Ti or pBR322 plasmids); or animal cell systems (Sambrook, supra; Ausubel et al., supra; Van Heeke, G. and S.M. Schuster (1989)).

Sci. USA 81:3655-3659; Harrington, J.J. et al. (1997) Nat. Genet. 15:345-355). Expression vectors derived from retroviruses, adenoviruses, or herpes or vaccinia viruses, or from various bacterial plasmids, may be used for delivery of polynucleotides to.

In bacterial systems, a number of cloning and expression vectors may be selected depending upon the use intended for polynucleotides encoding CGDD. For example, routine cloning, subcloning, and propagation of polynucleotides encoding CGDD can be achieved using a multifunctional E. coli vector such as PBLUESCRIPT (Stratagene, La Jolla CA) or pSPORT1 plasmid (Invitrogen). Ligation of polynucleotides encoding CGDD into the vector's multiple cloning site disrupts the lacZ gene, allowing a colorimetric screening procedure for identification of transformed bacteria containing recombinant molecules. In addition, these vectors may be useful for in vitro transcription, dideoxy sequencing, single strand rescue with helper phage, and creation of nested deletions in the.

264:5503-5509). When large quantities of CGDD are needed, e.g. for the production of antibodies, vectors which direct high level expression of CGDD may be used. For example, vectors containing the strong, inducible SP6 or T7 bacteriophage promoter may be used.

Yeast expression systems may be used for production of CGDD. A number of vectors containing constitutive or inducible promoters, such as alpha factor, alcohol oxidase, and PGH promoters, may be used in the yeast *Saccharomyces cerevisiae* or *Pichia pastoris*. In addition, such vectors direct either the secretion or intracellular retention of expressed proteins and enable integration of foreign polynucleotide sequences into the host genome.

complement of the polynucleotide encoding CGDD may be administered to a subject to treat or prevent a disorder associated. . . .

In other embodiments, any protein, agonist, antagonist, antibody, complementary sequence, or vector embodiment may be administered in combination with other appropriate therapeutic agents. . . .

Antisense sequences can also be introduced intracellularly through the use of viral vectors, such as retrovirus and adeno-associated virus vectors (Miller, A.D. (1990) Blood 76:27-31; Ausubel et al., supra; Uckert, W. and W. Walther (1994) Pharmacol. Ther. 63:323-347). . . . Other. . . .

In a further embodiment of the invention, diseases or disorders caused by deficiencies in CGDD are treated by constructing mammalian expression vectors encoding CGDD and introducing these vectors by mechanical means into CGDD-deficient cells. Mechanical transfer technologies for use with cells in vivo or ex vitro include (i) direct. . . .

Expression vectors that may be effective for the expression of CGDD include, but are not limited to, the pCDNA 3.1, EPITAG, PRCMV2, PREP, PVAX, PCR2-TOPOTA vectors (Invitrogen, Carlsbad CA), PCMV-SCRIPT, PCMV-TAG, PEGSH/PERV (Stratagene, La Jolla CA), and PTET-OFF, PTET-CN, PTRE2, PTRE2-LUC, PTK-HYG (Clontech, Palo Alto CA). CGDD may be. . . .

of the invention, diseases or disorders caused by genetic defects with respect to CGDD expression are treated by constructing a retrovirus vector consisting of (i) the . . . y

polynucleotide encoding CGDD under the control of an independent promoter or the retrovirus long terminal repeat (LTR). . . . RNA packaging signals, and (iii) a Rev-responsive element (RRE) along with additional retrovirus cis-acting RNA sequences and coding sequences required for efficient vector propagation. Retrovirus vectors (e.g., PB and PRENeo) are commercially available (Stratagene) and are based on published data (Riviere, I. et al. (1995) Proc. Natl. Acad. Sci. USA 92:6733-6737), incorporated by reference herein. The vector is propagated in an appropriate vector producing cell line (VPCL) that expresses an envelope gene with a tropism for receptors on the target cells or a promiscuous. . . .

transducing efficiency retroviral supernatant) discloses a method for obtaining retrovirus packaging cell lines and is hereby incorporated by reference. Propagation of retrovirus vectors, transduction of a population of cells (e.g., CD4<sup>+</sup> T-cells), and the return of transduced cells to a patient are procedures. . . .

cells which have one or more genetic abnormalities with respect to the expression of CGDD. The construction and packaging of

mammalian cells, a number of viral-based expression systems may be utilized. In cases where an adenovirus is used as an expression vector, polynucleotides encoding CGDD may be ligated into an adenovirus transcription/translation complex consisting of the late promoter and tripartite leader sequence. Insertion in. . . .

enhancers, such as the Rous sarcoma virus (RSV) enhancer, may be used to increase expression in mammalian host cells. SV40 or EBV-based vectors may also be used for high-level protein expression. . . .

stable expression of CGDD in cell lines is preferred. For example, polynucleotides encoding CGDD can be transformed into cell lines using expression vectors which may contain viral origins of replication and/or endogenous expression elements and a selectable marker gene on the same or on a separate vector. . . .

Following the introduction of the vector, cells may be allowed to grow for about 1 to 2 days in 74 enriched media before being switched to selective media. . . .

not only to identify transformants, but also to quantify the amount of transient or stable protein expression attributable to a specific vector system (Rhodes, C.A. (1995) Methods Mol. Biol. 55:121-131). . . .

Alternatively, polynucleotides encoding CGDD, or any fragments thereof, may be cloned into a vector for the production of an mRNA probe. Such vectors are known in the art, are commercially available, and may be used to synthesize RNA probes in vitro by addition of. . . .

from cell culture. The protein produced by a transformed cell may be secreted or retained intracellularly depending on the sequence and/or the vector used. As will be understood by those of skill in the art, expression vectors containing polynucleotides which encode CGDD may be designed to contain signal sequences which direct secretion of CGDD through a prokaryotic or eukaryotic. . . .

129/SvJ cell line, are derived from the early mouse embryo and grown in culture. The ES cells are transformed with a vector containing the gene of interest disrupted by a marker gene, e.g., the neomycin phosphotransferase gene (neo; Capecchi, M.R. (1989) Science 244:1288-1292). The vector integrates into the corresponding region of the host genome by homologous recombination. Alternatively, homologous recombination takes place using the Cre-loxP system to. . . .

In another embodiment, a vector capable of expressing CGDD or a fragment or derivative thereof may be administered to a subject to treat or prevent a. . . .

In an additional embodiment, a vector expressing the

adenovirus-based vectors are well known to those with ordinary skill in the art. Replication defective adenovirus vectors have proven to be versatile for importing genes encoding immunoregulatory proteins into intact islets in the pancreas (Csete, M.E. et al. (1995) Transplantation 27:263-268). Potentially useful adenoviral vectors are

described in U.S. Patent No. 5,707,618 to Armentano (Adenovirus vectors for gene therapy), hereby incorporated by reference. For adenoviral vectors, see also Antinozzi, P.A. et al. (1999; Annu.

which have one or more genetic abnormalities with respect to the expression of CGDD. The use of herpes simplex virus (HSV)-based vectors may be especially valuable for introducing CGDD to cells of the central nervous system, for which HSV has a tropism. The construction and packaging of herpes-based vectors are well known to those with ordinary skill in the art. A replication-competent herpes simplex virus (HSV) type I-based vector has been used to deliver a reporter gene to the eyes of primates (Liu, X. et al. (1999) Exp. Eye Res.

169:385-395). The construction of a HSV-1 virus vector has also been disclosed in detail in U.S.

For HSV vectors, see also Goins, W.F. et al. (1999; J. Virol. 73:519-532) and Xu, H. et al. (1994; Dev. Biol. 163:152-161). The manipulation.

In another embodiment, an alphavirus (positive, single-stranded RNA virus) vector is used to deliver polynucleotides encoding CGDD to target cells. The biology of the prototypic alphavirus, Semliki Forest Virus (SFV), has been studied extensively and gene transfer vectors have been based on the SFV genome (Garoff, H. and K.-J. Li (1998) Curr. Opin. Biotechnol. 9:464-469). During alphavirus RNA replication, a . . . region results in the production of a large number of CGDD-coding RNAs and the synthesis of high levels of CGDD in vector transduced cells. While alphavirus infection is typically associated with cell lysis within a few days, the ability to establish a persistent infection.

in vitro and in vivo transcription of DNA molecules encoding CGDD. Such DNA sequences may be incorporated into a wide variety of vectors with suitable RNA polymerase promoters such as T7 or SP6. Alternatively, these cDNA constructs that synthesize complementary RNA, constitutively or inducibly, can.

Many methods for introducing vectors into cells or tissues are available and equally suitable for use in vivo, in vitro, and ex vivo. For ex vivo therapy, vectors may be introduced into stem cells taken from the patient and clonally propagated for autologous transplant back into that same patient.

Means for producing specific hybridization probes for polynucleotides encoding CGDD include the cloning of polynucleotides encoding CGDD or CGDD derivatives

into vectors for the production of mRNA probes. Such vectors are known in the art, are commercially available, and may be used to synthesize RNA probes in vitro by means of.

was provided with RNA and constructed the corresponding cDNA libraries. Otherwise, cDNA was synthesized and cDNA libraries were constructed with the UNIZAP vector system (Stratagene) or SUPERScript plasmid system (Invitrogen), using the recommended procedures or similar methods known in the art (Ausubel et al. . . .

of cDNA Clones

Plasmids obtained as described in Example 1 were recovered from host cells by in vivo excision using the LTNIZAP vector system (Stratagene) or by cell lysis. Plasmids were purified using at least one of the following: a Magic or WIZARD Minipreps.

The polynucleotide sequences derived from Incyte cDNAs were validated by removing vector, linker, and poly(A) sequences and by masking ambiguous bases, using algorithms and programs based on BLAST, dynamic programming, and dinucleotide nearest.

plates, digested with CviJI cholera virus endonuclease (Molecular Biology Research, Madison Vt, and sonicated or sheared prior to religation into pUC 18 vector (Amersham Biosciences). For shotgun sequencing, the digested nucleotides were separated on low concentration

(0.6 to 0.8%) agarose gels, fragments were excised, and . . . agar digested with Agar ACE (Promega). Extended clones were religated using T4 ligase (New England Biolabs, Beverly MA) into pUC 18 vector (Amersham Biosciences), treated with Pfu DNA polymerase (Stratagene) to fill-in restriction site overhangs, and transfected into competent E. coli cells. Transformed cells. . . .

by requiring a minimum Phred quality score of 15, and removed sequence alignment errors and errors resulting from improper trinucleotide of vector sequences, chimeras, and splice variants. An automated procedure of advanced chromosome analysis analysed the original.

112 chromatogram files in the vicinity of the.

Microarray Preparation Sequences of the present invention are used to generate array elements. Each array element is amplified from bacterial cells containing vectors with cloned cDNA inserts. PCR amplification uses primers complementary to the vector sequences flanking the cDNA insert. Array elements are amplified in thirty cycles of PCR from an initial quantity of 1-2 ng.

CGDD is achieved using bacterial or virus-based expression systems. For expression of CGDD in bacteria, cDNA is subcloned into an appropriate vector containing an antibiotic resistance gene and an inducible promoter that directs high levels of cDNA

transcription. Examples of such promoters include, but . . . to, the trp-lac (tac) hybrid promoter and the T5 or T7 bacteriophage promoter in conjunction with the lac operator regulatory element. Recombinant vectors are transformed into suitable bacterial hosts, e.g., BL21(DE3).

by expressing the sequences encoding CGDD at physiologically elevated levels in mammalian cell culture systems. cDNA is subcloned into a mammalian expression vector containing a strong promoter that drives high levels of cDNA expression. Vectors of choice include PCMV SPORT plasmid (Invitrogen, Carlsbad CA) and PCR3.1 plasmid (Invitrogen), both of which contain the cytomegalovirus promoter. 5-10 /tg of recombinant vector are transiently transfected into a human cell line, for example, an endothelial or hematopoietic cell line, using either liposome formulations or electroporation. . . protein provides a means to distinguish transfected cells from nontransfected cells and is a reliable predictor of cDNA expression from the recombinant vector. Marker proteins of choice include, e.g., Green Fluorescent Protein (GFP; Clontech), CD64, or a CD64-GFP fusion protein. Flow cytometry (FCM), an. . .

progression when CGDD is expressed at physiologically elevated levels in mammalian cell culture systems. cDNA is subcloned into a mammalian expression vector containing a strong promoter that drives high levels of cDNA expression. Vectors of choice include PCMV SPORT (Life Technologies, Gaithersburg, IN) and PCR 3.1 (Invitrogen, Carlsbad, CA), both of which contain the cytomegalovirus promoter. 5-10 Mg of recombinant vector are transiently transfected into a human cell line, preferably of endothelial or hematopoietic origin, using either liposome formulations or electroporation. 1-2.1-tg. . . provides a means to distinguish transfected cells from nontransfected cells and is a reliable predictor of cDNA expression from the recombinant vector.

antisense CGDD RNA [Garkawsev, I. and K. Riabowol (1997) Mol. Cell Biol. 17:2014-2019]. cDNA encoding CGDD is subcloned into the pLNCX retroviral vector to enable expression of antisense CGDD RNA. The resulting construct is transfected into the ecotropic BOSC23 virus-packaging cell line. Virus contained. . .

Alternatively, CGDD can be expressed in a mammalian cell line by transfecting the cells with a eukaryotic expression vector encoding CGDD. Eukaryotic expression vectors are commercially available, and the techniques to introduce them into cells are well known to those skilled in the art. To assay. . .

can be measured by designing an antisense sequence to the 5' end of the gene and transfecting NIH 3T3 cells with a vector transcribing this sequence. The suppression of the endogenous gene will allow transformed fibroblasts to produce clumps of cells capable of forming metastatic. . .

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L3 ANSWER 1 OF 12  
PCTFULL, COPYRIGHT 2003 Univentio  
2003014322 PCTFULL, ED 20030303 EW 200308  
PROTEINS ASSOCIATED WITH CELL GROWTH, DIFFERENTIATION,  
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ET LA MORT CELLULAIRES  
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AGENT:  
LANGUAGE OF FILING:  
LANGUAGE OF PUBL.:  
DOCUMENT TYPE:  
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NUMBER KIND DATE  
WO 2003014322 A2 20030220

DESIGNATED STATES  
W:

AE AG AL AM AT AU AZ BA BB BG BR BY BZ CA CH CN CO CR  
CU CZ DE DK DM DZ EC EE ES FI GB GD GE GH GM HR HU ID  
IL IN IS JP KE KG KP KR KZ LC LK LR LS LT LU LV MA MD  
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AT BE CH CY DE DK ES FI FR GB GR IE IT LU MC NL PT SE  
TR

RW (ARIPO):  
RW (EAP):  
RW (EPO):  
RW (OAPI):

APPLICATION INFO.:  
PRIORITY INFO.:

BP BJ CF CG CI CM CA GN GQ GW ML MR NE SN TD TG  
US 2002-US25465 A 20020808  
US 2001-60/311,017 20010808  
US 2001-60/313,070 20010817  
US 2001-60/313,071 20010817  
US 2001-60/314,678 20010824  
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(FILE 'HOME' ENTERED AT 12:10:52 ON 20 MAR 2003)

FILE 'MEDLINE, CANCERLIT, BIOSIS, CONFSCI, SCISEARCH, CAPLUS, EMBASE,  
USPATFULL, PCTFULL' ENTERED AT 12:11:25 ON 20 MAR 2003

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L2 13 S LI AND VECTOR  
L3 12 DUP REM L2 (1 DUPLICATE REMOVED)

=> s l1 and DNA  
L4 24 L1 AND DNA

=> dup rem l4  
PROCESSING COMPLETED FOR L4  
L5 19 DUP REM L4 (5 DUPLICATES REMOVED)

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L5 ANSWER 1 OF 19 PCTFULL COPYRIGHT 2003 Univentio  
AN 2003014322 PCTFULL ED 20030303 EW 200308  
TIEN PROTEINES ASSOCIATED WITH CELL GROWTH, DIFFERENTIATION, AND DEATH  
TIFR PROTEINES ASSOCIEES A LA CROISSANCE, LA DIFFERENTIATION ET LA MORT  
CELLULAIRES  
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 WO 2003014322 A2 20030220  
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 RW (OAPI): BF BJ CF CG CI CM CA GN GQ GW ML MR NE SN TD TC  
 WO 2002-US25465 A 20020808  
 US 2001-60/311.017

NCL NCLM: 424/745.000  
 NCLS: 424/757.000; 424/756.000; 514/023.000; 514/027.000; 514/053.000;  
 424/755.000; 514/733.000  
 IC [7]  
 ICM: A61K035-78  
 ICS: A61K031-7C; A61K031-05  
 CAS INDEXING IS AVAILABLE FOR THIS PATENT.  
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 AN 2002:301167 USPTFULL  
 TI Nucleic acids, proteins, and antibodies  
 IN Rosen, Craig A.; Laytonville, MD, UNITED STATES  
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 CS School of Chemistry and Biochemistry, Georgia Institute of Technology,  
 Atlanta, GA 30332-0400, USA.  
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 CY United States  
 DT Journal: Article; (JOURNAL ARTICLE)  
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 Liska, DeAnn J.; Gig Harbor, WA, UNITED STATES  
 Tripp, Matthew, Gig Harbor, WA, UNITED STATES  
 Darland, Cary K.; Gig Harbor, WA, UNITED STATES  
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 DT Utility  
 FS APPLICATION  
 LN: CNT 2359  
 INCL INCLM: 424/745.000  
 INCLS: 424/757.000; 424/756.000; 514/023.000; 514/027.000; 514/053.000;  
 424/755.000; 514/733.000



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 95616, US  
 English  
 Patent  
 PI WO 2002029113 A2 20020411  
 W: AE AG AL AM AT AU AZ BA BB BG BR BY BZ CA CH CN CO CR CU CZ  
 DE DK DM DZ EC EE ES FI GB GD GE GH GM GR HU ID IL IN IS JP  
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 NO NZ PH PL PT RO RU SD SE SG SI SK SL TJ TM TR TT TZ UA UG  
 UZ VN YU ZA ZW  
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 TIEN PROTEINS RELATED TO SCHIZOPHRENIA AND USES THEREOF  
 TIFR PROTEINES LIEES A LA SCHIZOPHRENIE ET UTILISATIONS DE CELLES-CI  
 IN ST.GEORGE-HYSLOP, Peter, H., 210 Richview Avenue, Toronto, Ontario M5P  
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 RAE, Patricia, A., Sim & McBurney, 330 University Avenue, 6th Floor,  
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 W: AE AG AL AM AT AU AZ BA BB BG BR BY BZ CA CH CN CO CR CU CZ  
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 KE KG KP KR KZ LC LK LR LS LT LU LV MA MD MG MK MN MW MX MZ  
 NO NZ PH PL PT RO RU SD SE SG SI SK SL TJ TM TR TT TZ UA UG  
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 RW (ARIPO): GH GM KE LS MW MZ SD SL SZ TZ UG ZW  
 RW (EPO): AM AZ BY CY DE DK ES FI FR GB GR IE IT LU MC NL PT SE TR  
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 TIFR NOUVELLES PROTEINES ET ACIDES NUCLEIQUES LES CODANT  
 IN SPADERNA, Steven, K.;  
 TCHERNEV, Velizar;  
 LIU, Xiaohong;  
 SHENOY, Suresh;  
 SPYTEK, Kimberly;  
 ZERHUSEN, Bryan;  
 PATTURAJAN, Meera;  
 TAUPIER, Raymond, J.;  
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 GROSSE, William, M.;  
 SZEKERES, Edward, S.;  
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 LEPLEY, Denise, M.;

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DS	SHEN, Lei; BURGESS, Catherine, E.; SHIMKETS, Richard, A.; PADIGARU, Muralidhara CURAGEN CORPORATION; SPADERNA, Steven, K.; TCHERNEV, Velizar; LIU, Xiaohong; SHENOY, Suresh; SPYTEK, Kimberly; ZERHUSEN, Bryan; PATTURAJAN, Meera; TAUPIER, Raymond, J.; RASTELLI, Luca; GROSSE, William, M.; SZEKERES, Edward, S.; ALSOBROOK, John, II; LEPLEY, Denise, M.; SHEN, Lei; BURGESS, Catherine, E.; SHIMKETS, Richard, A.; PADIGARU, Muralidhara Patent WO 2002006339	AE AG AL AM AT AU AZ BA BB BG BR BY BZ CA CH CN CO CR CU CZ DE DK DM DZ EE ES FI GB GD GE GH GM GR HU ID IL IN IS JP KE KG KP KR KZ LC LK LR LS LT LU LV MA MD MG MK MN MW MX MZ NO NZ PL PT RO SD SE SG SI SK SL TJ TM TR TT TZ UA UG US UZ VN YU ZA ZW CH CM KE LS MW MZ SD SI SZ TZ UG ZW AM AZ BY BG KZ MD RU TJ TM AT BE CH CY DE DK ES FI FR GB GR IE IT LU MC NL PT SE TR BF BU CF CG CI CM GN GW ML MR NE SN TD TG	A2 20020124	AI PRAI	WO 2001-US2149 US 2000-60/215,854 US 2000-60/215,856 US 2000-60/215,902 US 2000-60/216,585 US 2000-60/216,586 US 2000-60/216,722 US 2000-60/218,622 US 2000-60/218,992 US 2000-60/221,285 US 2001-60/286,734 US 2001-60/274,260 US 2001-60/279,856 C07K014-47	PT SE TR BF BJ CF CG CI CM GN GW ML MR NE SN TD TG	WO 2001-US2149 US 2000-60/215,854 US 2000-60/215,856 US 2000-60/215,902 US 2000-60/216,585 US 2000-60/216,586 US 2000-60/216,722 US 2000-60/218,622 US 2000-60/218,992 US 2000-60/221,285 US 2001-60/286,734 US 2001-60/274,260 US 2001-60/279,856 C07K014-47	PT SE TR BF BJ CF CG CI CM GN GW ML MR NE SN TD TG
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US 2000-60/241.787	BANKAITIS-DAVIS, Danute, M.;		
US 2000-60/241.808	CHERONIS, John		
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US 2000-60/249.216	AU	Liu, Y. (1); wang, Z. (1); Kulesz-Martin, M. (1)	
US 2000-60/249.217	CS	(1) Dermatology, Oregon Health Sciences University, Portland, OR USA	
US 2000-60/249.211	SO	Journal of Investigative Dermatology, (August, 2001) Vol. 117, No. 2, pp. 475, print.	
US 2000-60/249.215		Meeting Info: 62nd Annual Meeting of the Society for Investigative	
US 2000-60/249.218		Dermatology Washington, DC, USA May 09-12, 2001	
US 2000-60/249.208	DT	Conference	
US 2000-60/249.213	LA	English	
US 2000-60/249.212	SL	English	
US 2000-60/249.207			
US 2000-60/249.245			
US 2000-60/249.244			
US 2000-60/249.297	L5	ANSWER 16 OF 19	PCTFULL
US 2000-60/249.214	AN	2000072854	PCTFULL
US 2000-60/249.219	TIEN	A DIETARY SUPPLEMENT CONTAINING VANADYL SULFATE, ALPHA-LIPOIC ACID, AND	
US 2000-60/249.264		TAURINE	
US 2000-60/249.209	TIFF	COMPLEMENT ALIMENTAIRE CONTENANT UN SULFATE VANADYL, UN ACIDE	
US 2000-60/249.300		ALPHA-LIPOIQUE ET UNE TAURINE	
US 2000-60/249.265	IN	PACIORETTY, Lindarp : SINDER, Stuart, J.	
US 2000-60/250.391	PA	BIONEXUS, LTD.;	
US 2000-60/256.719	LA	PACIORETTY, Linda	
US 2000-60/251.030	DT	Patent	
US 2000-60/251.988			



PI WO 2000072854 A1 20001207  
 DS AE AG AL AM AT AU AZ BA BB BG BR BY CA CH CN CR CU CZ DE DK  
 DM DZ EE ES FI GB GD GE GH GM HR HU ID IL IN IS JP KE KG KP  
 KR KZ LC LK LR LS LT LU LV MA MD MG MK MN MX MY NZ NO NZ PL  
 PT RO RU SD SE SG SI SK SL TJ TM TR TT TZ UA UG US UZ VN YU  
 ZA ZW GM KE LS MW MZ SD SL SZ T2 UG ZW AM AZ BY KG KZ MD  
 RU TJ TM AT BE CH CY DE DK ES FI FR GB GR IE IT LU MC NL PT  
 SE BF BJ CF CG CI CM GN GW ML MR NE SN TD TG  
 AI WO 2000-US15196 A 20000602  
 PRAI US 1999-60/137,080 A 19990602  
 L5 ANSWER 17 OF 19 PCTFULL COPYRIGHT 2003 Univentio  
 AN 200006069 PCTFULL ED 20020515  
 TIEN A PRESENILIN ASSOCIATED MEMBRANE PROTEIN AND USES THEREOF  
 TIFR PROTEINE MEMBRANAIRE ASSOCIEE A LA PRESENILINE ET SES UTILISATIONS  
 IN ST. GEORGE-HYSLOP, Peter, H.;  
 FRASER, Paul, E.  
 PA THE GOVERNING COUNCIL OF THE UNIVERSITY OF TORONTO  
 LA English  
 DT Patent  
 DS WO 2000060069 A1 20001012  
 W: AE AG AL AM AT AU AZ BA BB BG BR BY CA CH CN CR CU CZ DE DK  
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 AT BE CH CY DE DK ES FI FR GB GR IE IT LU MC NL PT SE BF BJ  
 CF CG CI CM CA CN GM ML MR NE SN TD TG  
 AI WO 2000-CA354 A 20000403  
 PRAI US 1999-60/127,452 19990401  
 ICM US 1999-60/173,826 19991230  
 CS C07K014-705; A01K067-027; C12N005-10; C120001-68; G01N033-50  
 L5 ANSWER 18 OF 19 MEDLINE  
 AN 2000020628 MEDLINE  
 DN 20200628 Pubmed ID: 10734318  
 TI SPAP, a new AAA-protein specific to early spermatogenesis and  
 malignant conversion.  
 AU Liu Y; Black J; Kiesel N; Kulesz-Martin M F  
 CS Program of Biochemistry and Department of Pharmacology and Therapeutics,  
 Roswell Park Cancer Institute, Elm & Carlton Streets, Buffalo, New York,  
 NY 14263, USA.  
 NC CA16056 (NCI)  
 SO ONCOGENE, (2000 Mar 16) 19 (12) 1579-88.  
 CY Journal code: 8711562. ISSN: 0950-9232.  
 DT ENGLAND: United Kingdom  
 LA Journal: Article; (JOURNAL ARTICLE)  
 FS English  
 OS Priority Journals  
 EM GENBANK-AF049099  
 ED 200004  
 Entered STN: 20000505  
 Last Updated on STN: 20000505  
 Entered Medline: 20000421  
 L5 ANSWER 19 OF 19 MEDLINE  
 AN 94368094 MEDLINE  
 DN 94368094 Pubmed ID: 8085823  
 TI Genes involved in self-protection against the lantibiotic subtilin  
 produced by Bacillus subtilis ATCC 6633.  
 AU Klein C; Enian K D  
 CS Institute for Microbiology, Biozentrum der Johann Wolfgang  
 Goethe-Universitat, Frankfurt am Main, Federal Republic of Germany.

SO APPLIED AND ENVIRONMENTAL MICROBIOLOGY, (1994 Aug) 60 (8) 2793-801.  
 CY Journal code: 7605801. ISSN: 0099-2240.  
 DT United States  
 LA Journal: Article; (JOURNAL ARTICLE)  
 FS English  
 OS Priority Journals  
 EM GENBANK-U09819  
 ED 199410  
 Entered STN: 19941021  
 Last Updated on STN: 19941021  
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 L6 13 L1 AND CDNA  
 => dup rem l6  
 PROCESSING COMPLETED FOR L6  
 L7 12 DUP REM L6 (1 DUPLICATE REMOVED)  
 => d 1-12  
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 TIEN PROTEINS ASSOCIATED WITH CELL GROWTH, DIFFERENTIATION, AND DEATH  
 TIFR PROTEINES ASSOCIEES A LA CROISSANCE, LA DIFFERENTIATION ET LA MORT  
 CELLULAIRES  
 IN AZIMZAI, Valda, 5518 Boulder Canyon Drive, Castro Valley, CA 94552, US  
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for US only;  
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for US only;  
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for US only;  
XU, Yuming, 1739 Walnut Drive, Mountain View, CA 94040, US [US, US];  
for US only;  
YAO, Monique, G., 1189 Woodgate Drive, Carmel, IN 46033, US [US, US];  
for US only;  
YUE, Henry, 826 Lois Avenue, Sunnyvale, CA 94087, US [US, US];  
for US only;  
YUE, Huibin, 1170 South Stelling Road, Cupertino, CA 95014, US [US, US];  
for US only;  
HAMLET-COX, Diana, Incyte Genomics, Inc., 3160 Porter Drive, Palo Alto, CA 94304, US [US, US];  
English  
LAF Patent  
DT English  
FI WO 20030014322 A2 20030220  
DS W: AE AG AL AM AT AU AZ BA BB BG BR BY BZ CA CH CN CO CR CU CZ DE DK DM E2 EC EE ES FI GB GD GE GH GM HR HU ID IL IN IS JP KE KG KP KR LC LK LR LS LT LU LV MA MD MG MK MN MW MX MZ NO NZ OM PH PL PT RO RU SD SE SG SI SK SL TJ TM TN TR TT TZ UA US UZ VN YU ZA ZW ZW  
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RW (EAP): AM AZ BY KG KZ MD RU TJ TM  
RW (EPO): AT BE CH CY DE DK ES FI FR GB GR IE IT LJ MC NL PT SE TR  
RW (OAPI): BF BJ CF CG CI CM GA GN GQ GW ML MR NE SN TD TG  
AI WO 2002-US25465 A 20020808  
PRAI US 2001-60/311.017 20010808  
US 2001-60/313.070 20010817  
US 2001-60/313.071 20010817  
US 2001-60/314.678 20010824  
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US 2001-60/317.913 20010907  
US 2001-60/322.182 20010914  
US 2001-60/340.747 20011207  
US 2001-60/342.761 20011220  
US 2002-60/369.129 20020329  
L7 ANSWER 2 OF 12 CAPLUS COPYRIGHT 2003 ACS DUPLICATE 1  
AN 2002:755245 CAPLUS  
DN 137:274175  
TI DNA, cDNA and protein sequences of spermatogenesis assocd.  
IN Kulesz-Martin, Molly F.; Liu, Yuangang  
PA USA Pat. Appl. Publ., 42 pp., Cont.-in-part of U.S. Ser. No. 777,753.  
SO CODEN: USXXCO  
DT Patent  
LA English  
FAN CNT 1  
PI US 2002143169 A1 20021003 APPLICATION NO. DATE  
US 2001-850697 20010508

PRAI US 1997-938308 B2 19970927  
US 2001-777753 A2 20010206

L7 ANSWER 3 OF 12 USPATFULL  
AN 2002:301167 USPATFULL  
TI Nucleic acids, proteins, and antibodies  
IN Ruben, Craig A., Laytonville, MD, UNITED STATES  
Barash, Steven C., Rockville, MD, UNITED STATES  
US 2002168711 A1 20021114  
AI 2001-764868 A1 20010117 (9)  
US 2000-179065P 20000131 (60)  
PRAI US 2000-180628P 20000204 (60)  
US 2000-214886P 20000628 (60)  
US 2000-217487P 20000711 (60)  
US 2000-225758P 20000814 (60)  
US 2000-220963P 20000726 (60)  
US 2000-217496P 20000711 (60)  
US 2000-225447P 20000814 (60)  
US 2000-218290P 20000714 (60)  
US 2000-225757P 20000814 (60)  
US 2000-226868P 20000822 (60)  
US 2000-216647P 20000707 (60)  
US 2000-225267P 20000814 (60)  
US 2000-216880P 20000707 (60)  
US 2000-225270P 20000814 (60)  
US 2000-251869P 20001208 (60)  
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US 2000-234274P 20000921 (60)  
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US 2000-241809P 20001020 (60)  
US 2000-249299P 20001117 (60)  
US 2000-236327P 20000929 (60)  
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US 2000-244617P 20001101 (60)  
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US 2000-236370P 20000929 (60)  
US 2000-236802P 20001002 (60)  
US 2000-237037P 20001002 (60)  
US 2000-237040P 20001002 (60)  
US 2000-240960P 20001020 (60)  
US 2000-239935P 20001013 (60)  
DT Utility  
FS APPLICATION  
LN.CNT 31967  
INCL INCLM: 435/069.100

INCLM: 435/325.000; 435/320.100; 435/183.000; 530/350.000; 536/023.100  
NCLM: 435/069.100  
NCLS: 435/325.000; 435/320.100; 435/183.000; 530/350.000; 536/023.100  
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ICM: C12P021-02  
ICS: C12N005-06; C07H021-04; C12N009-00; C07K014-435  
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L7 ANSWER 4 OF 12 USPATFULL  
AN 2002:112541 USPATFULL  
TI Proteins related to schizophrenia and uses thereof  
IN St. George-Hyslop, Peter H., Toronto, CANADA  
Fraser, Paul E., Toronto, CANADA  
PA The Governing Council of the University of Toronto (non-U.S. corporation)  
PI US 2002058276 A1 20020516  
AI US 2001-945258 A1 20010831 (9)  
PRAI US 2000-229889P 20000901 (60)  
DT Utility  
FS APPLICATION  
LN.CNT 2909  
INCL INCLM: 435/006.000  
INCLM: 424/009.200; 803/003.000  
NCLM: 435/006.000  
NCLS: 424/009.200; 803/003.000  
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ICM: C12Q001-68  
ICS: A61K049-00; A01K057-00  
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L7 ANSWER 5 OF 12 PCTFULL COPYRIGHT 2003 Univentio  
AN 200209062 PCTFULL ED 20021218 EW 200250  
TIEN NOVEL ANTIBODIES THAT BIND TO ANTIGENIC POLYPEPTIDES, NUCLEIC ACIDS  
ENCODING THE ANTIGENS, AND METHODS OF USE  
TIFR NOUVEAUX ANTICORPS SE FIXANT A DES POLYPEPTIDES ANTIGENIQUES, ACIDES  
NUCLEIQUES CODANT LES ANTIGENES ET MODES D'UTILISATION  
IN ANDERSON, David, W., 85 Montoya Drive, Branford, CT 06405, US [US, US];  
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ELARIFI, Ivor R. Mintz, Levin, Cohn, Ferris, Glovsky, and Popeo, P.,	
C., One Financial Center, Boston, MA 02111, US	
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ANSWER 2002046385	PCTFULL
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TIER	
IN	
TANG, Y., Tom, 4230 Ranwick Court, San Jose, CA 95118, US [US, US]:	
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 HILLMAN, Jennifer, L., 230 Monroe Drive, #17, Mountain View, CA 94040,  
 US [US, US], for US only;  
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PA

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 LA English  
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 US 2001-60/262,839 20010119  
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 L7 ANSWER 7 OF 12 PCTFULL COPYRIGHT 2003 Univentio  
 AN 2002029113 PCTFULL ED 20020627 EW 200215  
 TIEN METHODS FOR MONITORING MULTIPLE GENE EXPRESSION  
 TIFR METHODES DE SURVEILLANCE DE L'EXPRESSION GENETIQUE MULTIPLE  
 IN BERKA, Randy, 3609 Modoc, Davis, CA 95616, US;  
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 AG STARNES, Robert, Novozymes Biotech, Inc., 1445 Drew Avenue, Davis, CA  
 95616, US  
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 US 2001-60/279,526 20010327  
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 L7 ANSWER 8 OF 12 PCTFULL COPYRIGHT 2003 Univentio  
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 TIEN PROTEINS RELATED TO SCHIZOPHRENIA AND USES THEREOF  
 TIFR PROTEINES LIEES A LA SCHIZOPHRENIE ET UTILISATIONS DE CELLES-CI  
 IN ST.GEORGE-HVSLOP, Peter, H., 210 Richview Avenue, Toronto, Ontario M5P  
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 FRASER, Paul, E., 611 Windermere Avenue, Toronto, Ontario M6S 3L9, CA  
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 AG RAE, Patricia, A., Sim & McBurney, 330 University Avenue, 6th Floor,  
 Toronto, Ontario M5G 1R7, CA  
 LAF English  
 LA English

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L7 ANSWER 9 OF 12 PCTFULL COPYRIGHT 2003 Univentio  
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TIEN NOVEL PROTEINS AND NUCLEIC ACIDS ENCODING SAME  
TIFF NOUVELLES PROTEINES ET ACIDES NUCLEIQUES LES CODANT  
IN SPADERNA, Steven, K.;  
TCHERNEV, Velizar;  
LIU, Xiaohong;  
SHENOY, Suresh;  
SPYTEK, Kimberly;  
ZERHUSEN, Bryan;  
PATTURAJAN, Meera;  
TAUPIER, Raymond, J.;  
RASTELLI, Luca;  
GROSSE, William, M.;  
SZEXERES, Edward, S.;  
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SHEN, Lei;  
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SPADERNA, Steven, K.;  
TCHERNEV, Velizar;  
LIU, Xiaohong;  
SHENOY, Suresh;  
SPYTEK, Kimberly;  
ZERHUSEN, Bryan;  
PATTURAJAN, Meera;  
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ALSOBROOK, John, II;  
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L7 ANSWER 10 OF 12 PCTFULL COPYRIGHT 2003 Univentio  
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TIFF ACIDES NUCLEIQUES, PROTEINES ET ANTICORPS  
IN ROSEN, Craig, A.;  
BARASH, Steven, C.;  
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HUMAN GENOME SCIENCES, INC.;  
ROSEN, Craig, A.;  
BARASH, Steven, C.;  
RUBEN, Steven, M.  
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WO 2001-CA1243 A 20010831  
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ICM C07K014-47

L7 ANSWER 9 OF 12 PCTFULL COPYRIGHT 2003 Univentio  
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BURGESS, Catherine, E.;  
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L7 ANSWER 10 OF 12 PCTFULL COPYRIGHT 2003 Univentio  
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RUBEN, Steven, M.;  
HUMAN GENOME SCIENCES, INC.;  
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ICM AG1K048-00  
ICS C12N005-00

L7 ANSWER 11 OF 12 PCTFULL COPYRIGHT 2003 Univentio  
AN 2000060069 PCTFULL ED 20020515  
TIEN A PRESENILIN ASSOCIATED MEMBRANE PROTEIN AND USES THEREOF  
TIFFR PROTEINE MEMBRANAIRE ASSOCIEE A LA PRESENILINE ET SES UTILISATIONS  
IN ST. GEORGE-HYSLOP, Peter, H.;

FRASER, Paul, E.  
THE GOVERNING COUNCIL OF THE UNIVERSITY OF TORONTO  
English  
Patent  
PI WO 2000060069  
DS W: AE AG AL AM AT AU AZ BA BB BG BR BY CA CH CN CR CU CZ DE DK  
DM DZ EE ES FI GB GD GE GH GM HR HU ID IL IN IS JP KE KG KP  
KR KZ LC LK LR LS LT LU LV MA MD MG MK MN MW MX NZ PL PT  
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GH GM KE LS MW SD SL SZ TZ UG ZW AM AZ BY KG KZ MD RU TJ TM  
AT BE CH CY DE DK ES FI FR GB GR IE IT LU MC NL PT SE BF BJ  
CF CG CI CM GN GW ML MR NE SN TD TG

AI WO 2000-CA354  
PRAI US 1999-60/127,452 19990401  
US 1999-60/173,826 19991230  
ICM C12N015-12  
ICS C07K014-705; A01K067-027; C12N005-10; C12Q001-68; G01N033-50

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L7 ANSWER 12 OF 12 CAPLUS COPYRIGHT 2003 ACS
AN 2000:242308 CAPLUS
DN 133:29064
TI SPAP, a new AAA-protein specific to early spermatogenesis and
   malignant conversion
AU Liu, Yungang, Black, Jennifer; Kisiel, Nicholas; Kulesz-Martin, Molly F.
CS Program of Biochemistry and Department of Pharmacology and Therapeutics,
   Roswell Park Cancer Institute, Buffalo, NY, 14263, USA
SO Oncogene (2000), 19(12), 1579-1588
   CODEN: ONCNE5; ISSN: 0950-9232
PB Nature Publishing Group
DT Journal
LA English
RE.CNT 44 THERE ARE 44 CITED REFERENCES AVAILABLE FOR THIS RECORD
        ALL CITATIONS AVAILABLE IN THE RE FORMAT

```

```

=> d his

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(FILE 'HOME' ENTERED AT 12:10:52 ON 20 MAR 2003)
FILE 'MEDLINE, CANCERLIT, BIOSIS, CONFSCI, SCISEARCH, CAPLUS, EMBASE,
USPATFULL, PCTFULL' ENTERED AT 12:11:25 ON 20 MAR 2003
L1 202 S SPAP
L2 13 S L1 AND VECTOR
L3 12 DUP REM L2 (1 DUPLICATE REMOVED)
L4 24 S L1 AND DNA
L5 19 DUP REM L4 (5 DUPLICATES REMOVED)
L6 13 S L1 AND CDNA
L7 12 DUP REM L6 (1 DUPLICATE REMOVED)

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```

=> s atcc (a) 98558
L8 0 ATCC (A) 98558

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```

=>

```

```

---Logging off of STN---

```

```

=>
Executing the logoff script...

```

```

=> LOG Y

```

COST IN U.S. DOLLARS	SINCE FILE	TOTAL
FULL ESTIMATED COST	ENTRY	SESSION
	60.59	60.80

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STN INTERNATIONAL LOGOFF AT 12:23:53 ON 20 MAR 2003

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